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(54) Ink-jet recording method

Tintenstrahlzeichnungsverfahren
Méthode d'enregistrement à jet d'encre

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- DATABASE WPI Week 8926 Derwent Publications Ltd., London, GB; AN 89-209986 XP002012612 & JP-A-01 146 784 (FUJI PHOTO FILM)
- DATABASE WPI Week 8929 Derwent Publications Ltd., London, GB; AN 89-209987 XP002012613 & JP-A-01 146 785 (FUJI PHOTO FILM)
- DATABASE WPI Week 8848 Derwent Publications Ltd., London, GB; AN 88-341382 XP002012614 & JP-A-63 252 779 (CANON)

Remarks:

The file contains technical information submitted after the application was filed and not included in this specification

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Description

[0001] The present invention relates to a recording method for ink-jet recording, and particularly to a recording method for ink-jet recording in which a water based ink is used.

[0002] Recently, following the proliferation of personal computers, ink-jet recording system printers are rapidly spreading. Specifically, in the field of graphic art and designing, their utilization is being taken notice of due to their high quality resembling photography.

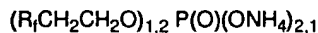
[0003] As a recording sheet used for an ink-jet recording system, conventionally, a recording sheet wherein an ink receiving layer (hereinafter, referred also to as "ink-absorption layer") is provided on ordinary paper or a support (referred to as an ink-jet recording paper) has been used. However, when the above-mentioned recording papers are employed, much ink blurring results and glossiness is low. Therefore, the above-mentioned recording papers could not be employed for the above-mentioned field wherein high resolution and high glossiness is required.

[0004] In addition, when a transparent support is used to produce an original for an OHP (over-head projector), there was the problem that a porous ink-absorption layer interfered light transmittance.

[0005] In order to overcome the above-mentioned problems, Japanese Patent Publication Open to Public Inspection (hereinafter, referred to as Japanese Patent O.P.I. Publication) Nos. 216990/1992 and 64306/1994 disclose technologies of an ink-jet recording sheet wherein a resin-covered paper, i.e., an RC (resin coated) paper, in which both sides of the paper are covered with resin, is used for a support and gelatin is used for the ink-receiving layer.

[0006] According to the above-mentioned specifications, a high resolution and a glossy image could be obtained. However, they were still inadequate in terms of overall quality. Specifically, dot reproducibility was insufficient so that they could not be used for original graphic art work nor for printing of a fine design picture.

[0007] US 4,781,985 discloses an ink jet recording transparency which comprises a transparent resin base which is coated with a cationic fluorine-containing surfactant or with an anionic fluorine-containing surfactant of the formula:



wherein R_f is fluoroalkyl having from 3 to 8 carbons.

[0008] An objective of the present invention is to provide an ink-jet recording method using a recording sheet for ink-jet recording capable of obtaining high image quality and stable image quality.

[0009] The above objects of the invention can be attained by an ink-jet recording method using the following:

1. A recording sheet for ink-jet recording comprising a support, and provided thereon, an ink receiving layer, wherein the ink receiving layer contains an anionic fluorine-containing surfactant and a cationic fluorine-containing surfactant.

2. The recording sheet for ink-jet recording of 1 above, wherein the content ratio of the anionic fluorine-containing surfactant to the cationic fluorine-containing surfactant is 1:10 to 10:1 in terms of mole ratio.

3. The recording sheet for ink-jet recording of 1 or 2 above, wherein the content ratio of the anionic fluorine-containing surfactant to the cationic fluorine-containing surfactant is 3:7 to 7:3 in terms of mole ratio.

4. The recording sheet for ink-jet recording of 1, 2 or 3 above, wherein the ink receiving layer further contains gelatin.

5. The recording sheet for ink-jet recording of 1, 2, 3 or 4 above, wherein the ink receiving layer further contains gelatin and a water-soluble polymer.

6. The recording sheet for ink-jet recording of 1, 2, 3, 4 or 5 above, wherein the water-soluble polymer is at least one compound selected from polyvinyl pyrrolidones and polyvinyl alcohols.

7. The recording sheet for ink-jet recording of 1, 2, 3, 4, 5 or 6 above, wherein the content ratio of the water-soluble polymer to the gelatin is 10 to 70 weight %.

8. The recording sheet for ink-jet recording of 1, 2, 3, 4, 5, 6 or 7 above, wherein the content ratio of the water-soluble polymer to the gelatin is 40 to 60 weight %.

9. The recording sheet for ink-jet recording of 1, 2, 3, 4, 5, 6, 7 or 8 above, wherein the support is a resin-covered paper, in which both surfaces of the paper are covered with a resin, or a polyester film.

10. The recording sheet for ink-jet recording of 9 above, wherein the resin is a polyolefin resin.

11. The recording sheet for ink-jet recording of 9 above, wherein the polyester is polyethylene terephthalate.

The invention provides the following:

12. A method for ink-jet recording, the method employing a recording sheet for ink-jet recording comprising an ink receiving layer provided on a support, wherein the ink receiving layer comprises an anionic fluorine-containing surfactant and a cationic fluorine-containing surfactant.

The surfactants are defined below.

13. The method for ink-jet recording of 12 above, wherein the content ratio of the anionic fluorine-containing sur-

factant to the cationic fluorine-containing surfactant is from 1:10 to 10:1 in terms of mole ratio.

14. The method for ink-jet recording of 12 or 13 above, wherein the content ratio of the anionic fluorine-containing surfactant to the cationic fluorine-containing surfactant is from 3:7 to 7:3 in terms of mole ratio.

15. The method for ink-jet recording of 12, 13 or 14 above, wherein the ink receiving layer further comprises gelatin.

5 16. The method for ink-jet recording of claim 12, 13, 14 or 15 above, wherein the ink receiving layer further contains gelatin and a water-soluble polymer.

17. The method for ink-jet recording of 12, 13, 14, 15 or 16 above, wherein the water-soluble polymer is at least one compound chosen from polyvinyl pyrrolidones and polyvinyl alcohols.

10 18. The method for ink-jet recording of 12, 13, 14, 15, 16 or 17 above, wherein the content ratio of the water-soluble polymer to the gelatin is from 10 to 70 weight %.

19. The method for ink-jet recording of 12, 13, 14, 15, 16, 17 or 18 above, wherein the content ratio of the water-soluble polymer to the gelatin is from 40 to 60 weight %.

20. The method for ink-jet recording of 12, 13, 14, 15, 16, 17, 18 or 19 above, wherein the support is a resin covered paper, in which both surfaces of the paper are covered with a resin, or a polyester film.

15 21. The method for ink-jet recording of 20 above, wherein the resin is a polyolefin resin.

22. The method for ink-jet recording of 20 above, wherein the polyester is polyethylene terephthalate.

[0010] The present invention will be detailed below:

20 [0011] The recording sheet for use in the ink-jet recording method of the invention is characterized in that the ink receiving layer comprises anionic fluorine-containing surfactant and cationic fluorine-containing surfactant, as defined below. It has been so far considered that use of a combination of an anionic surfactant and a cationic surfactant causes undesirable coagulation in a coating solution prior to coating. However, experimental results from the present inventors have shown that a fluorine-containing surfactant does not cause coagulation in solution condition. In addition, when the fluorine-containing surfactant is used for recording sheets for ink-jet recording, excellent ink-receivability results and the phenomenon whereby an ink spot is difficult to diffuse over the passage of time, scarcely occurs. Therefore, larger ink spots can be utilized and it has been found that an image with high density and little unevenness can be obtained.

25 [0012] The anionic fluorine-containing surfactant is chosen from $C_{10}F_{21}(CH_2)_{10}COOH$ and those represented by the following Formula (FA):

30 (FA) $R_f-(D)_t-Y$ Formula

wherein R_f represents a fluorine-containing alkyl group having from 2 to 30 carbon atoms or an aryl group having a fluorine-containing alkyl group of from 2 to 30 carbon atoms; D represents a divalent group having at least one of -O-, -COO-, -CON(R_1)- and -SO₂N(R_1)- linking groups and having from 1 to 12 carbon atoms; R_1 represents an alkyl group having from 1 to 5 carbon atoms; t represents 1 or 2; Y represents -COOM-, -SO₃M or -OSO₃M; and M represents a hydrogen atom, an alkaline metal or a quaternary ammonium salt.

35 [0013] R_f preferably represents a fluorine-containing alkyl group having from 3 to 10 carbon atoms, D preferably represents a divalent group containing an alkylene group having from 1 to 5 carbon atoms and at least one of -O-, -COO-, -CON(R_1)- and -SO₂N(R_1)-, or a divalent group containing an arylene group and at least one of -O-, -COO-, -CON(R_1)- and -SO₂N(R_1)- in which R_1 represents an alkyl group having from 1 to 5 carbon atoms, and Y preferably represents -COOM or -SO₃M in which M represents an alkali metal, preferably sodium or potassium.

40 [0014] Next, practical examples of the anionic surfactants will be illustrated. However, the present invention is not limited thereto.

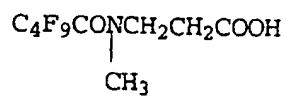
45 FA-2 $C_{10}F_{21}(CH_2)_{10}COOH$

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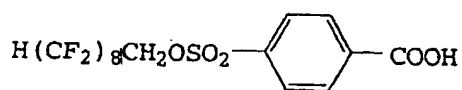
$$C_8F_{17}SO_2N \begin{array}{c} | \\ C_3H_7 \end{array} CH_2CH_2O(CH_2)_3SO_3Na$$

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FA-11

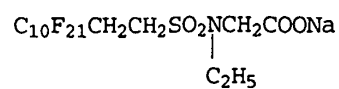


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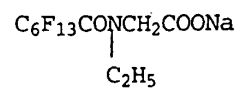
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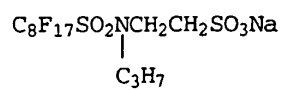
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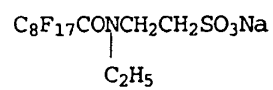
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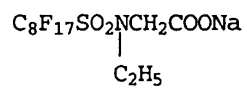
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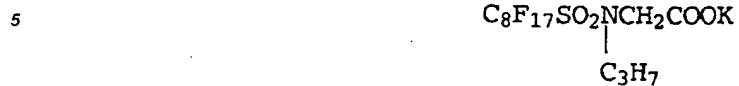
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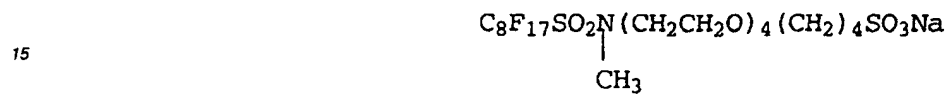
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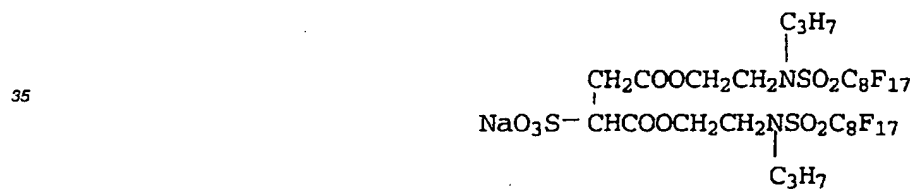


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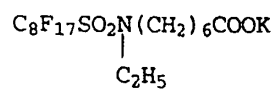
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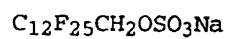
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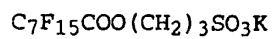
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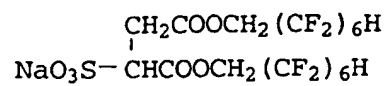
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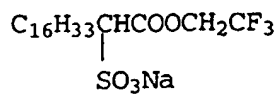
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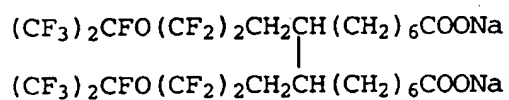
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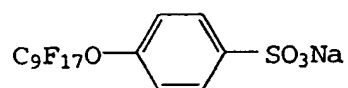
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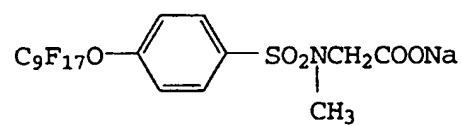
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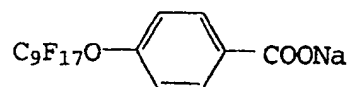
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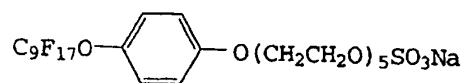


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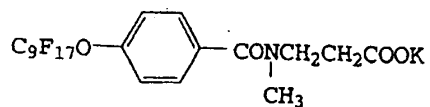
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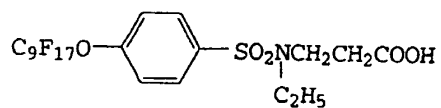
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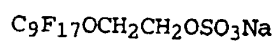
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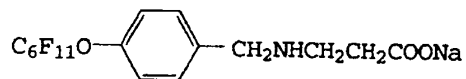


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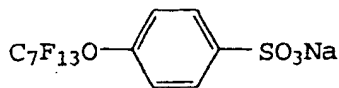
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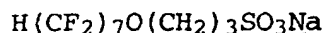
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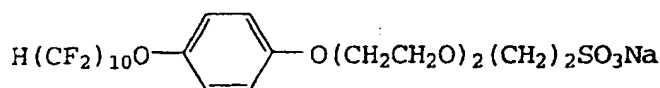
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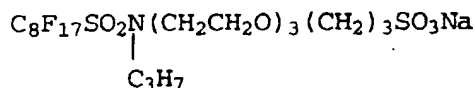
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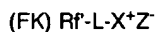


FA-49



[0015] Specifically preferable are anionic fluorine-containing surfactants comprising at least one $-\text{SO}_2\text{N}(\text{R}')-$ bond.

[0016] Cationic fluorine-containing surfactants used in the present invention are compounds represented by the following Formula (FK):



Formula

wherein R' represents a hydrocarbon group having from 1 to 20 carbon atoms in which at least one hydrogen atom is substituted by a fluorine atom; L represents a chemical bond or a divalent group; X represents a cation; and Z represents a counter anion.

[0017] Examples of R' include $-\text{C}_k\text{F}_{k+1}$ ($k=1$ to 20, preferably 3 to 12), $-\text{C}_m\text{F}_{2m-1}$ ($m=2$ to 20, preferably 3 to 12).

[0018] Examples of L include $-\text{SO}_2\text{N}(\text{R}')(\text{CH}_2)_p-$, $-\text{CON}(\text{R}')(\text{CH}_2)_p-\text{OASO}_2\text{N}(\text{R}')(\text{CH}_2)_p-$, $-\text{OACON}(\text{R}')(\text{CH}_2)_p-$, $-\text{OAO}(\text{CH}_2)_p-$, $-\text{OA}(\text{CH}_2)_p-$, $-\text{O}(\text{CH}_2\text{CH}_2\text{O})_q(\text{CH}_2)_p-$, $-\text{O}(\text{CH}_2)_p-$, $-\text{N}(\text{R}')(\text{CH}_2)_p-$, $-\text{SO}_2\text{N}(\text{R}')(\text{CH}_2)_p\text{O}(\text{CH}_2)_r-$, $-\text{CON}(\text{R}')(\text{CH}_2)_p\text{O}(\text{CH}_2)_r-$, $-\text{OASO}_2\text{N}(\text{R}')(\text{CHR}')_p\text{OA}-$ and $-(\text{CH}_2)_p(\text{CHOH})_s(\text{CH}_2)_r-$, in which A represents alkylene or arylene.

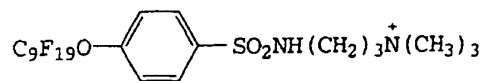
[0019] Examples of X^+ include $-\text{N}^+(\text{R}')_3$, $-\text{N}^+(\text{CH}_2\text{CH}_2\text{OCH}_3)_3$, $-\text{N}^+\text{C}_4\text{H}_8\text{O}(\text{R}')$, $-\text{N}^+(\text{R}')(\text{R}^2)(\text{CH}_2\text{CH}_2\text{OCH}_3)$, $-\text{N}^+\text{C}_5\text{H}_5$, $-\text{N}^+(\text{R}')(\text{R}^2)(\text{CH}_2)_p\text{C}_6\text{H}_5$ and $-\text{N}^+(\text{R}')(\text{R}^2)(\text{R}^2)$, wherein R' and R^2 independently represent a hydrogen atom or an alkyl group, which may have a substituent, having from 1 to 6 carbon atoms; p , r and s independently represent from 0 to 6; and q represents from 1 to 20.

[0020] Examples of Y include I^- , Cl^- , Br^- , CH_3SO_3^- and $\text{CH}_3-\text{C}_6\text{H}_4-\text{SO}_3^-$.

[0021] Particular examples of the cationic fluorine-containing surfactants preferably used in the present invention are set out below. However, the present invention is not limited thereto.

FK-1

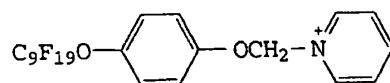
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 I^-

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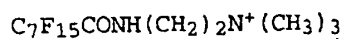
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 Cl^-

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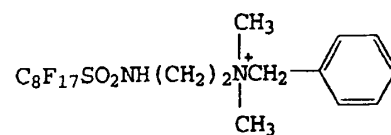
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 Cl^-

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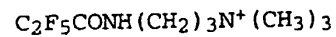
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 Cl^-

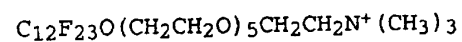
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FK-5

 I^-

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FK-6

 I^-

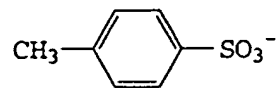
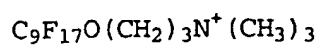
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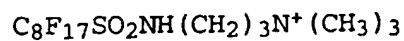
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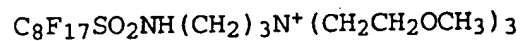
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FK-8.

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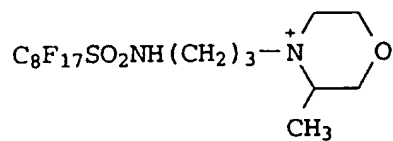
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FK-10

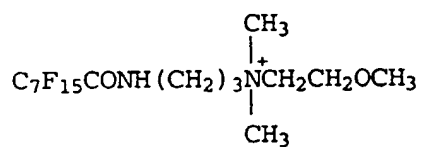


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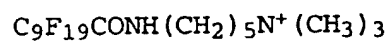
FK-11

Br⁺

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FK-12



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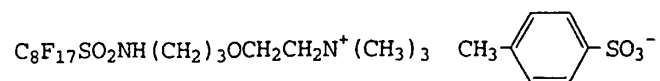
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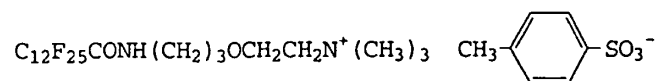
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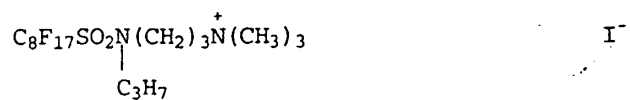
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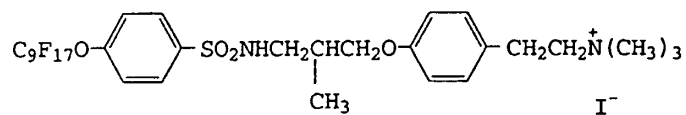
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FK-16

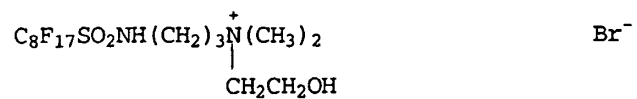
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FK-17

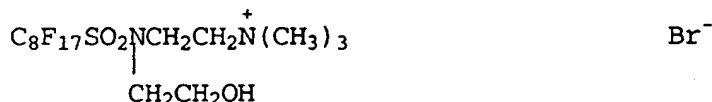
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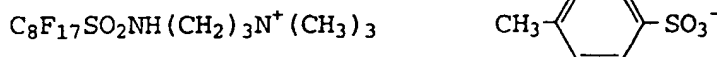
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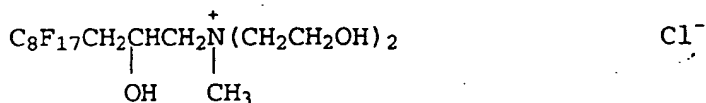
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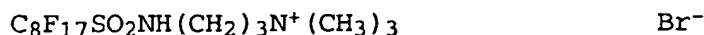
FK-19



FK-20



FK-21



[0022] The anionic fluorine-containing surfactants or the cationic fluorine-containing surfactants of the present invention can be synthesized by methods described in U.S. Patent Nos. 2,559,751, 2,567,011, 2,732,398, 2,764,602, 2,806,866, 2,809,998, 2,915,376, 2,915,528, 2,918,501, 2,934,450, 2,937,098, 2,957,031, 3,472,894 and 3,555,089, British Patent Nos. 1,143,927 and 1,130,822, Japanese Patent Publication No. 37304/1970, Japanese Patent O.P.I. Nos. 9613/1972, 134614/1974, 117705/1975, 117727/1975, 121243/1975, 41182/1977 and 12392/1976, J. Chem. Soc., 1950, page 2789 and 1957, pp. 2574 and 2640, J. Amer. Chem. Soc., Volume 79, page 2549 (1957), J. Japan Oil Chemists Soc., Volume 12, page 653 and J. Org. Chem., Volume 30, page 3524 (1965).

[0023] Some of the above-mentioned fluorine-containing surfactants are commercially available as follows: Megafac F produced by DaiNippon Ink Chemical Industry Co. Ltd.; Fluorad FC produced by Minesota Mining and Manufacturing Company; Monflor produced by Imperial Chemical Industry; Zonyls produced by E. I. Du Pont Nemeras and Company; Licowet produced by Falbewereke Hexist.

[0024] The total amount of the anionic fluorine-containing surfactant and cationic fluorine-containing surfactant used in the invention is preferably from 0.1 to 1000 mg, more preferably from 0.5 to 300 mg and further more preferably from 1.0 to 150 mg per 1 m² of recording sheet. The anionic fluorine-containing surfactant and cationic fluorine-containing surfactant each may be used in two or more kinds, respectively. In addition to these surfactants, a nonionic or amphoteric fluorine-containing surfactant or a hydrocarbon type surfactant may be used in combination.

[0025] The addition amount ratio of the anionic fluorine-containing surfactant to the cationic fluorine-containing surfactant in the invention is preferably from 1:10 to 10:1 by mole ratio, and more preferably from 3:7 to 7:3 by mole ratio.

[0026] In the invention, the ink receiving layer of a recording sheet for ink-jet recording comprises the anionic fluorine-containing surfactant and the cationic fluorine-containing surfactant, and optionally further comprises a binder for forming the layer. The binder may include a natural polymer and a synthetic polymer. The binder used in the invention is preferably gelatin.

[0027] As gelatin, any gelatin made from animal collagen can be used, but gelatin made from pig skin, cow skin or cow bone collagen is preferable. The kind of gelatin is not specifically limited, but lime-processed gelatin, acid proc-

essed gelatin or gelatin derivatives (for example, gelatin derivatives disclosed in Japanese Patent Publication Nos. 38-4854/1962, 39-5514/1964, 40-12237/1965, 42-26345/1967 and 2-13595/1990, U.S. Patent Nos. 2,525,753, 2,594,293, 2,614,928, 2,763,639, 3,118,766, 3,132,945, 3,186,846 and 3,312,553 and British Patent Nos. 861,414 and 103,189) can be used alone or in combination.

5 [0028] In one of the preferable embodiments of the invention, acid processed gelatin is used in view of ink absorption, and gelatin in which the amino group is inactivated is used in view of ink absorption or glossiness.

[0029] The acid processed gelatin preferably used in the invention is prepared by deliming collagen and then processing it with an acid such as hydrochloric acid.

10 [0030] The amino group inactivating gelatin preferably used in the invention is gelatin in which 50% or more, preferably 80% or more and more preferably 90%, of the amino groups, are inactivated. The above gelatin, which is prepared according to a conventional acetylation method, includes acetylated gelatin, phthaloylated gelatin, malenoylated gelatin, benzoylated gelatin, succinoylated gelatin, methyl urea gelatin, phenylcarbamoylated gelatin, and carboxy modified gelatin.

15 [0031] The jelly strength of gelatin used in the invention is preferably not less than 150 kg, and more preferably from 200 to 300 kg (according to the PAGI method). The jelly strength of gelatin is measured with a bloom gelometer.

[0032] In the invention, the gelatin content of the ink receiving layer is preferably from 3 to 50 g/m², and more preferably from 5 to 30 g/m². If the gelatin content is less than 3 g/m², ink reception is deteriorated, resulting in ink release from the ink received portions after printing, and if the gelatin content exceeds 50 g/m², ink reception is improved but cracking or curling occurs.

20 [0033] The thickness of the ink receiving layer in the invention is preferably from 1 to 50 μ m, and more preferably from 2 to 30 μ m.

[0034] In order to improve ink reception or dot reproduction, the following water-soluble polymer can be used. The water-soluble polymer includes polyvinyl alcohol, polyvinyl pyrrolidone, polyvinyl pyridinium halide, modified polyvinyl alcohol such as polyvinyl formal or their derivatives (see Japanese Patent O.P.I. Publication Nos. 145879/1985, 25 220750/1985, 143177/1986, 235182/1986, 235183/1986, 237681/1986 and 261089/1986), an acryl group-containing polymer such as polyacrylamide, polydimethylacrylamide, polydimethylaminoacrylate, polysodiumacrylate, acrylic acid-meth acrylic acid copolymer salt, polysodiummethacrylate or acrylic acid-vinyl alcohol copolymer (disclosed in Japanese Patent O.P.I. Publication Nos. 168651/1985 and 9988/1987), a natural polymer or its derivatives such as starch, oxidation starch, carboxylated starch, dialdehyde starch, cationated starch, dextrin, sodium alginate, gum arabic, casein, pullulan, dextrane, methylcellulose, ethylcellulose, carboxymethylcellulose or hydroxypropylcellulose (Japanese Patent O.P.I. Publication Nos. 174382/1974, 262685/1985, 143177/1986, 181679/1986, 193879/1986 and 287782/1986), a synthetic polymer such as polyethylene glycol, polypropylene glycol, polyvinyl ether, polyglycerin, maleic acid-alkylvinylether copolymer, maleic acid-N-vinylpyrrole copolymer, styrene-maleic anhydride copolymer or polyethylene imine (disclosed in Japanese Patent O.P.I. Publication Nos. 32787/1986, 237680/1986 and 277483/1986). 30 Of these, polyethylene glycol, polyvinyl alcohol and polyvinyl pyrrolidone are preferred and polyvinyl alcohol and polyvinyl pyrrolidone are particularly preferred. The average molecular weight of these polymers is preferably not less than 10,000, and more preferably not less than 100,000 in view of tackiness of the polymer surface.

[0035] The water soluble polymer can be used alone or in combination, but it is preferred that the water soluble polymer is used in combination with the above described gelatin. The water soluble polymer content of the ink receiving 40 layer is from 3 to 50 g/m², and preferably from 5 to 30 g/m².

[0036] When the water soluble polymer is used with gelatin, the addition amount ratio (by weight %) of the water soluble polymer to gelatin depends on the type of polymer, but is preferably from 10 to 70%, more preferably from 40 to 60% by weight in order to improve ink reception and/or dot image reproduction.

45 [0037] The ink receiving layer preferably further contains basic latexes disclosed in Japanese Patent Application No. 7-54286/1995, and especially preferably exemplified compounds I-1 to I-12 on pages 16 to 18, in that excellent print can be obtained independent of circumstances.

[0038] The ink receiving layer preferably contains water soluble polymers disclosed in Japanese Patent Application No. 7-64334/1995 in order to shorten the ink drying time.

50 [0039] In the material used in the method of invention, the ink receiving layer can be hardened with a hardener in order to improve water resistance or dot reproduction. Examples of the hardener include aldehyde compounds such as formaldehyde and glutaraldehyde, ketone compounds such as diacetyl and chloropentanedion, bis(2-chloroethylurea), 2-hydroxy-4,6-dichloro-1,3,5-triazine, reactive halogen-containing compounds disclosed US Patent No. 3,288,775, divinylsulfone, reactive olefin-containing compounds disclosed US Patent No. 3,635,718, N-methylol compounds disclosed US Patent No. 2,732,316, isocyanates disclosed US Patent No. 3,103,437, aziridine compounds disclosed US Patent 55 Nos. 3,017,280 and 2,983,611, carbodiimides disclosed US Patent No. 3,100,704, epoxy compounds disclosed US Patent No. 3,091,537, a halogencarboxyaldehyde such as mucochloric acid, a dioxane derivative such as dihydroxy dioxane, and inorganic hardeners such as chromium alum, potash alum and zirconium sulfate. These hardeners can be used alone or in combination. The addition amount of hardener is preferably 0.01 to 10g, and more preferably 0.1 to

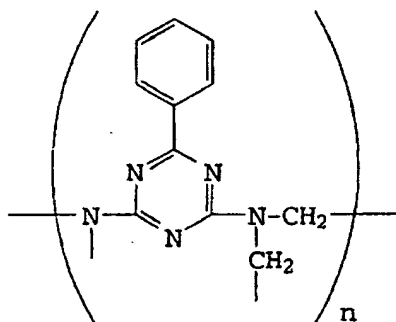
5 g based on 100g of a binder contained in the ink receiving layer.

[0040] The ink receiving layer may contain a matting agent in an amount of 0.005 to 0.1 g/m² in order to prevent adhesion defect such as blocking.

[0041] The matting agent can be defined as discontinuously dispersed particles such as inorganic or organic materials capable of being dispersed in a hydrophilic organic colloid. The inorganic matting agent includes oxides such as silicon oxide, titanium oxide, magnesium oxide and aluminum oxide, alkali earth metal salts such as barium sulfate, calcium carbonate, and magnesium sulfate, light-insensitive silver halide particles such as silver chloride and silver bromide (each of which may contain a small amount of an iodine atom), and glass. Besides these substances are used inorganic matting agents which are disclosed in West German Patent No. 2,529,321, British Patent Nos. 760 775 and 1,260,772, US Patent Nos. 1,201,905, 2,192,241, 3,053,662, 3,062,649, 3,257,296, 3,322,555, 3,353,958, 3,370,951, 3,411,907, 3,437,484, 3,523,022, 3,615,554, 3,635,714, 3,769,020, 4,021,245 and 4,029,504.

[0042] The organic matting agent includes starch, cellulose ester such as cellulose acetate propionate, cellulose ether such as ethyl cellulose and a synthetic resin. The synthetic resin is a water insoluble or sparingly soluble polymer which includes a polymer of an alkyl(meth)acrylate, an alkoxyalkyl(meth)acrylate, a glycidyl(meth)acrylate, a (meth)acrylamide, a vinyl ester such as vinyl acetate, acrylonitrile, an olefin such as ethylene, or styrene and a copolymer of the above described monomer with other monomers such as acrylic acid, methacrylic acid, α,β -unsaturated dicarboxylic acid, hydroxyalkyl(meth)acrylate, sulfoalkyl(meth)acrylate and styrene sulfonic acid. Further, a benzoguanamin-formaldehyde resin, an epoxy resin, nylon, polycarbonates, phenol resins, polyvinyl carbazol or polyvinylidene chloride can be used. Besides the above are used inorganic matting agents which are disclosed in British Patent No. 1,055,713, US Patent Nos. 1,939,213, 2,221,873, 2,268,662, 2,322,037, 2,376,005, 2,391,181, 2,701,245, 2,992,101, 3,079,257, 3,262,782, 3,443,946, 3,516,832, 3,539,344, 554, 3,591,379, 3,754,924 and 3,767,448, Japanese Patent O.P.I. Publication Nos. 49-106821/1974 and 57-14835/1982.

[0043] Of these, preferred are polymethylmethacrylate, a benzoguanamine-formaldehyde polycondensate (a benzoguanamine resin as represented by the following formula, for example, Eposter produced by Nihon Shokubai Kagaku-kogyo Co., Ltd., or (Chemical Substance Registry No. 7-31 compound),



polyolefins (for example, Frobeads LE-1080, CL-2080, HE-5023: produced by Seitetsu Kagaku Co., Ltd., or Chemipar V-100 produced by Mitsui Sekiyu Kagaku Kogyo Co., Ltd.), polystyrene beads (produced by Moritex Co., Ltd.), nylon beads (produced by Moritex Co., Ltd.), AS resin beads (produced by Moritex Co., Ltd.), epoxy resin beads (produced by Moritex Co., Ltd.) or polycarbonate resin beads (produced by Moritex Co., Ltd.).

[0044] These matting agents may be used in combination.

[0045] The ink receiving layer may further comprise, in addition to the above surfactants the binder and the hardener, various conventional additives such as inorganic pigment, colorants, colored pigment, a fixing agent for ink dyes, a UV absorber, an anti-oxidant, a dispersing agent, an anti-foaming agent, a leveling agent, an antiseptic agent, a brightening agent, a viscosity stabilizing agent and a pH adjusting agent.

[0046] As a coating method of an ink receiving layer coating solution, any conventional coating method (for example, a curtain method, an extrusion method, an air-knife method, a roll coating method and a rod bar coating method) can be used.

[0047] The ink receiving layer may be single layered or multi-layered. Examples of multi-layered structures include those disclosed in Japanese Patent O.P.I. Publication Nos. 57-89954/1982, 60-224578/1985 and 61-12388/1988. Further, an ink penetrating layer may be provided on the ink receiving layer.

[0048] The ink receiving layer is provided on at least one surface of the support, and may be provided on both surfaces of the support in order to minimize curling.

[0049] As the support, a transparent or opaque support can be optionally used according to its use. A conventional

support can be used as the transparent support, which includes a film or plate of polyester resins, cellulose acetate resins, acryl resins, polycarbonate resins, polyvinyl chloride resins, polyimide resins, cellophane or celluloid and a glass plate.

[0050] The thickness of the transparent support is preferably from 10 to 200 μm . As the opaque support, any conventional one such as paper, coat paper, synthetic paper, resin-covered paper, pigment-containing opaque film or foaming film can be used in the invention, but synthetic paper, a resin-covered paper or various films are preferable in view of glossiness or smoothness, and resin-covered paper or polyester film is preferable in view of touch or luxuriousness.

[0051] The base paper constituting the resin-covered paper used in the method of the invention is not specifically limited, and any conventional paper can be used, but a smooth paper used as a conventional photographic support is preferable. As pulp constituting the base paper, natural pulp, reproduction pulp or synthetic pulp is used singly or in admixture. These base papers may contain additives such as a sizing agent, a reinforcing agent, a filler, an anti-static agent, a fluorescent brightening agent or a dye which is usually used in paper manufacture. A surface sizing agent, a surface reinforcing agent, a fluorescent brightening agent, an antistatic agent and an anchoring agent may be coated on the surface of the material.

[0052] The thickness of the base paper is not specifically limited, but is preferably from 10 to 200 μm . A base paper having a smooth surface is preferable, which is obtained by applying pressure to or calendering, paper, during or after papering. The weight of the base paper is preferably from 30 to 250 g/m^2 .

[0053] As a resin for a resin-covered paper, a polyolefin resin or a resin capable of being hardened with an electron beam can be used. The polyolefin resin includes an olefin homopolymer such as a low density polyethylene, a high density polyethylene, polypropylene or polypentene, an olefin copolymer such as ethylene-propylene copolymer or their mixture, each having various densities or melt viscosity indexes (melt index). These resins can be used singly or in combination.

[0054] The resin for the resin-covered paper preferably contains various additives, for example, white pigment such as titanium oxide, zinc oxide, talc or calcium carbonate, a fatty acid amide such as stearic acid amide or arachidic acid amide, a fatty acid metal salt such as zinc stearate, calcium stearate, aluminum stearate or magnesium stearate, an anti-oxidant such as Irganox 1010 or Irganox 1076, blue pigment or dyes such as cobalt blue, ultramarine, or phthalocyanine blue, magenta pigment or dyes such as cobalt violet, fast violet or manganese violet, a brightening agent and a UV absorber. These additives can be suitably used in combination.

[0055] The resin-covered paper, which is the support preferably used in the method of the invention, is manufactured by a so-called extrusion method casting a thermally fused resin (for example, fused polyolefin) on the moving paper, whereby both surfaces of the paper are covered with the resin. When the paper is covered with a resin capable of being hardened with electron beam irradiation, the resin is coated with a conventional coater such as a gravure coater or a blade coater and then is irradiated with electron beam to harden the coated resin. Before the paper is coated with a resin, the surface of the paper is preferably subjected to activation treatment such as corona discharge treatment or flame treatment. The surface of the support on the ink receiving layer side is glossy or matte depending upon its usage, and a glossy surface is preferable. The back side of a support is not necessarily covered with a resin, but is preferably covered with a resin in view of prevention of curling. The back surface of a support is ordinarily non-glossy, but the back surface or both surfaces of the support are optionally subjected to activation treatment such as corona discharge treatment or flame treatment. The thickness of a covered resin is not specifically limited, but is ordinarily from 5 to 50 μm .

[0056] The water based ink herein referred to is a recording liquid comprising the following colorants, solvents and other additives. The colorant includes a direct dye, an acid dye, a basic dye, a reactive dye and food dyes.

[0057] The solvent for the water based ink includes alkyl alcohols having from 1 to 4 carbon atoms such as methyl alcohol, ethyl alcohol, isopropyl alcohol, butyl alcohol, sec-butyl alcohol, tert-butyl alcohol and iso-butyl alcohol, amides such as dimethylformamide and dimethylacetamide, ketones or ketonealcohols such as acetone and diacetone alcohol, ethers such as tetrahydrofuran and dioxane, polyalkylene glycols such as polyethylene glycol and polypropylene glycol, alkylene glycols having from 2 to 6 carbon atoms such as ethylene glycol, propylene, butylene glycol, triethylene glycol, 1,3,6-hexane triol, hexylene glycol, thiodiglycol and diethylene glycol, polyhydric alcohol lower alkyl ethers such as glycerin, ethylene glycol methylether, diethylene glycol methyl(or ethyl)ether and triethylene glycol monomethylether, pyrrolidinones such as 2H-pyrrolidinone, and pyrrolidones such as 1-methyl-2-pyrrolidone and 2-pyrrolidone. Of these water soluble solvents, a polyhydric alcohol such as diethylene polyhydric alcohol lower alkyl ethers such as triethylene glycol monomethylether and triethylene glycol monoethylether, and pyrrolidones are preferable.

[0058] The solvent for the ink is preferably a mixture of water and the above described organic solvent in view of prevention of ink head nozzle clogging. The mixture ratio of water to the organic solvent is preferably from 1:9 to 9:1 by weight, and more preferably from 4:6 to 9:1 by weight.

[0059] The additives include a pH adjusting agent, a metal chelating agent, an anti-fungal, a viscosity adjusting agent, a surface tension adjusting agent, a wetting agent, a surfactant and an anti-rust agent.

EXAMPLES

[0060] The invention will be detailed in the following examples, but the invention is not limited thereto.

Exempl 1

[0061] The following ink receiving layer coating solution was coated on RC paper available on the market by means of a bar coating method and dried to give a dry thickness of 8 g/m². Thus, a recording sheet sample for ink jet recording was obtained. Next, yellow (Y), magenta (M) and cyan (C) color images were printed on the above obtained sample, employing an ink jet printer, Design Jet 650C (produced by Hewlett Packard Co., Ltd.). The resulting sample was evaluated for dot diameter, optical density and unevenness at magenta color image portions.

(Ink receiving layer aqueous coating solution)

[0062]

| | | |
|---|------------------------------|---------------------|
| Lime-processed gelatin (Bloom strength; 250 g) | | 100 parts by weight |
| Surfactants (shown in Table 1) | | shown in Table 1 |
| * Solid component concentration of the coating solution was 8% by weight. The composition of each ink was as follows: | | |
| Y: | Direct yellow 50 (Cl. 29025) | 6 parts by weight |
| | Diethylene glycol | 47 parts by weight |
| | Water | 47 parts by weight |
| M: | Xylene red B (Cl. 45100) | 6 parts by weight |
| | Diethylene glycol | 47 parts by weight |
| | Water | 47 parts by weight |
| C: | Light green SF yellowish | 6 parts by weight |
| | Diethylene glycol | 47 parts by weight |
| | Water | 47 parts by weight |

[0063] Results of evaluation will be shown as follows:

(Evaluation of image unevenness)

[0064]

- A: Excellent, (without problem).
- B: Good.
- C: Some patchiness observed.
- D: Patchiness prevalent too much to be practically used.

Table 1

| Sample No. | Anionic fluorine-containing surfactant (mg/m ²) | Cationic fluorine-containing surfactant (mg/m ²) | Dot diameter (μm) | Density of solid image | Image unevenness | Remarks |
|------------|---|--|-------------------|------------------------|------------------|---------|
| 1-1 | None | None | 100 | 1.01 | D | Comp. |
| 1-2 | FA-2 (4) | None | 102 | 1.02 | C | Comp. |
| 1-3 | None | FK-4 (4) | 103 | 1.04 | C | Comp. |
| 1-4 | FA-2 (4) | FK-4 (4) | 102 | 1.21 | B | Inv. |
| 1-5 | FA-16 (3) | FK-5 (3) | 103 | 1.22 | B | Inv. |
| 1-6 | FA-18 (4) | FK-8 (4) | 100 | 1.23 | B | Inv. |
| 1-7 | FA-19 (4) | FK-21 (4) | 104 | 1.21 | A | Inv. |
| 1-8 | FA-31 (4) | FK-20 (4) | 101 | 1.21 | B | Inv. |
| 1-9 | FA-2 (4) | FK-4 (4) | 101 | 1.18 | B | Inv. |
| 1-10 | FA-2 (5.6) | FK-4 (4) | 101 | 1.20 | B | Inv. |
| 1-11 | FA-2 (2.4) | FK-4 (4) | 102 | 1.20 | B | Inv. |
| 1-12 | FA-2 (0.4) | FK-4 (4.0) | 101 | 1.17 | B | Inv. |

Comp. : Comparative

Inv. : Invention

[0066] From the above Table 1, it can be seen that the recording sheet for ink jet recording of the present invention could obtain favorable results in all evaluation items. Therefore, it is evident that the recording sheet of the present invention can produce images with high quality.

Example 2

[0067] Recording sheets for ink-jet recording were prepared in the same manner as in Sample 1-6 prepared in Example 1, except that the amount of the surfactants was changed as shown in the following Table 2. The resulting sheets were evaluated in the same manner as in Example 1, except that dot diameter, optical density and image unevenness at cyan color image portions were evaluated. In addition, glossiness of the print image was evaluated visually.

(Evaluation of glossiness)

[0068]

A: Excellent

B: Glossiness of the printed portions is slightly degraded.

C: Glossiness of the printed portions is totally absent.

[0069] Table 2 shows the results.

Table 2

| Sample No. | Anionic surfactant (mg/m ²) | Cationic surfactant (mg/m ²) | Optical density | Image unevenness | Glossiness |
|------------|---|--|-----------------|------------------|------------|
| 2-1 | 500 | 500 | 1.22 | B | C |
| 2-2 | 300 | 300 | 1.23 | A | C |
| 2-3 | 150 | 150 | 1.24 | A | B |

Table 2 (continued)

| Sample No. | Anionic surfactant (mg/m ²) | Cationic surfactant (mg/m ²) | Optical density | Image unevenness | Glossiness |
|------------|--|---|-----------------|------------------|------------|
| 2-4 | 100 | 100 | 1.21 | A | B |
| 2-5 | 75 | 75 | 1.21 | A | A |
| 2-6 | 25 | 25 | 1.20 | A | A |
| 2-7 | 5 | 5 | 1.19 | A | A |
| 2-8 | 0.5 | 0.5 | 1.17 | A | A |
| 2-9 | 0.25 | 0.25 | 1.17 | B | A |
| 2-10 | 0.05 | 0.05 | 1.15 | C | A |

[0070] From the above-mentioned Table 2, it can be understood that the recording sheet for ink-jet recording of the present invention has no unevenness at an image portion and in addition, has excellent glossiness.

Example 3

[0071] Recording sheets for ink jet recording were prepared in the same manner as in Sample 1-6 of Example 1, except that water-soluble polymers described in Table 3 were further added. They were evaluated for ink absorption in addition to the same evaluation as described in Example 1.

(Evaluation on ink absorptivity)

[0072] Commercially available wood free paper was brought into contact with the magenta solid image at 1 kg/m² pressure, and then, the transfer degree of the ink onto the wood free paper was evaluated.

A : When the contact was carried out 30 seconds after printing, no ink transfer was observed.

B : When the contact was carried out 30 seconds after printing, slight ink transfer was observed, but when the contact was carried out one minute after printing, no ink transfer was observed.

C : When the contact was carried out one minute after printing, slight ink transfer was observed, but no practical problem.

Table 3

| Sample No. | Water-soluble polymer, addition ratio based on gelatin (wt%) | Optical density of solid image | Image unevenness | Ink absorptivity |
|------------|--|--------------------------------|------------------|------------------|
| 3-1 | PEG 6000 ¹⁾ (50%) (Nippon Yushi) | 1.34 | B | C |
| 3-2 | Cerazol 100A ²⁾ (50%) (Meisei Kagaku) | 1.44 | B | A |
| 3-3 | PVA GL-05 ³⁾ (50%) (Nihon Gosei Kagaku Kogyo) | 1.32 | A | B |
| 3-4 | PVP K-90 ⁴⁾ (5%) (BASF) | 1.25 | B | C |
| 3-5 | PVP K-90 (10%) | 1.28 | B | B |
| 3-6 | PVP K-90 (30%) | 1.30 | B | A |
| 3-7 | PVP K-90 (40%) | 1.33 | A | A |
| 3-8 | PVP K-90 (50%) | 1.36 | A | A |
| 3-9 | PVP K-90 (60%) | 1.37 | A | A |
| 3-10 | PVP K-90 (70%) | 1.33 | B | A |
| 3-11 | PVP K-90 (80%) | 1.34 | B | B |

¹⁾ average molecular weight: 6,000

²⁾ average molecular weight: 150,000

³⁾ average molecular weight: 22,000

⁴⁾ average molecular weight: 360,000

[0073] As is apparent from the results shown in Table 3, when the water-soluble polymer is added to the ink-receiving layer, both the optical density of the print portion and the absorptivity of the ink are improved.

Example 4

[0074] A sample was prepared with the same coating solution composition as Example 1 except that a commercially available polyethylene terephthalate film (the layer thickness was 100 μ m) in place of the RC paper used in Example 1, and was evaluated in the same manner as in Example 1. The results were almost the same as Example 1.

Example 5

[0075] Samples 5-1 and 5-2 were prepared in the same manner as in Sample 3-6 of Example 3 except that cationic denatured PVP (GAFQUAT HS-100 produced by ISP Inc.) and polyallyl amine hydrochloride (PAA-HCl produced by Nit-toh-bo Co. Ltd.) were added in an amount of 10 wt% based on gelatin, respectively, and were evaluated in the same manner as in Example 3. Results showed that Samples 5-1 and 5-2 had the same optical density and uneven image as those of Sample 3-6. However, the ink absorption of Samples 5-1 and 5-2 were half of that of Sample 3-6. Therefore, noticeable improvement was observed.

Example 6

[0076] The 7 weight % (as solid content) aqueous solutions containing the following layer compositions were coated on RC paper according to a bar coating method to obtain a recording sheet for ink jet recording.

| | | |
|---|--|----------------------|
| First Layer (a layer closest to the polyethylene paper) | | |
| Lime-processed gelatin | | 1.2 g/m ² |
| PVP-K90 | | 0.8 g/m ² |
| Second Layer (an intermediate layer) | | |
| Gelatin (as shown in Table 4) | | 3.3 g/m ² |
| PVP-K90 | | 3.7 g/m ² |
| Third Layer (a layer furthest from the polyethylene paper) | | |
| Gelatin (as shown in Table 4) | | 0.7 g/m ² |
| PVP-K90 | | 0.6 g/m ² |
| Sodium di-2-ethylhexylsulfosuccinate | | 16 mg/m ² |
| Exemplified Compound FA-19 | | 5 mg/m ² |
| Exemplified Compound FA-21 | | 5 mg/m ² |
| Matting agent (Polymethylmethacrylate average grain size : 10 μm) | | 40 mg/m ² |

[0077] An image was printed using the above obtained sheet and an ink-jet printer (MJ-700V2C produced by Epson Co., Ltd.).

[0078] The resulting sheet was evaluated for image unevenness and ink absorption in the same manner as in Example 3. Further, glossiness of non-image portions at 60° was measured by a gloss meter VG-ID (produced by Nihon Den-syoku kogyo Co., Ltd.). The evaluation criteria of the glossiness were as follows:

- A : 80 or more; excellent glossiness
- B : Less than 80 to 70; good glossiness
- C : Less than 70 to 50; fair glossiness
- D : Less than 50; poor glossiness

[0079] The results are shown in Table 4.

Table 4

| Sample No. | Gelatin in Second Layer | Gelatin in Third Layer | Image unevenness | Ink absorptivity | Glossiness at non-image portions |
|------------|-------------------------|------------------------|------------------|------------------|----------------------------------|
| 6-1 | lime-processed gelatin | lime-processed gelatin | B | B | B |
| 6-2 | * PC gelatin | lime-processed gelatin | A | B | B |
| 6-3 | lime-processed gelatin | PC gelatin | B | A | A |
| 6-4 | PC gelatin | PC gelatin | A | A | A |
| 6-5 | acid-processed gelatin | lime-processed gelatin | B | A | B |

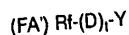
* PC gelatin: phenylcarbamoylated gelatin a phenylcarbamoylation degree of 98% or more

[0080] As is apparent from Table 4, use of acid-processed gelatin provides more excellent ink absorptivity, and the use of amino group inactivated gelatin provides more excellent glossiness of non-image portions as well as providing more excellent image quality and ink absorptivity, resulting in improvement of print quality.

Claims

1. A method for ink-jet recording, comprising the step of:

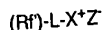
jetting a water-based ink onto a recording sheet, the recording sheet comprising a support and provided therein, an ink receiving layer comprising a binder, an anionic fluorine-containing surfactant and a cationic fluorine-containing surfactant, wherein the anionic fluorine-containing surfactant is $C_{10}F_{21}(CH_2)_{10}COOH$ or a compound of the following formula (FA):



wherein R_f represents a fluorine-containing alkyl group having from 2 to 30 carbon atoms or an aryl group having a fluorine-containing alkyl group of from 2 to 30 carbon atoms; D represents a divalent group having at least one of $-O-$, $-COO-$, $-CON(R_1)-$ and $-SO_2N(R_1)-$ linking groups and having from 1 to 5 carbon atoms; R_1 represents an alkyl group having from 1 to 5 carbon atoms; t represents 1 or 2; Y represents $-COOM-$, $-SO_3M$ or OSO_3M ; and M represents a hydrogen atom an alkaline metal or a quaternary ammonium salt.

2. A method according to claim 1, wherein D is a divalent group having an $-SO_2N(R_1)-$ bond and R_1 is an alkyl group having from 1 to 5 carbon atoms.

3. A method according to claim 1 or 2, wherein the cationic fluorine-containing surfactant is a compound of the following formula (FK):



wherein Rf' represents a fluorine-containing hydrocarbon group having from 1 to 20 carbon atoms; L represents a chemical bond or $-SO_2N(R^1)(CH_2)_p-$, $-CON(R^1)(CH_2)_p-$, $-OASO_2N(R^1)(CH_2)_p-$, $-OACON(R^1)(CH_2)_p-$, $-OAO(CH_2)_p-$, $-OA(CH_2)_p-$, $-O(CH_2CH_2O)_q(CH_2)_p-$, $-O(CH_2)_p-$, $-N(R^1)(CH_2)_p-$, $-SO_2N(R^1)(CH_2)_pO(CH_2)_r-$, $-CON(R^1)(CH_2)_pO(CH_2)_r-$, $-OASO_2N(R^1)(CHR^1)_pOA-$ or $-(CH_2)_p(CHOH)_s(CH_2)_r-$, in which A represents an alkylene group or an arylene group, each R^1 and R^2 independently represent a hydrogen atom or an alkyl group having from 1 to 6 carbon atoms, p , r and s independently represent from 0 to 6 and q represents from 1 to 20; X^+ represents a cation; and Z^- represents an anion.

4. A method according to claim 3, wherein L is chosen from $-SO_2N(R^1)(CH_2)_p-$, $-CON(R^1)(CH_2)_p-$, $-OASO_2N(R^1)(CH_2)_p-$, $-OACON(R^1)(CH_2)_p-$, $-OAO(CH_2)_p-$, $-OA(CH_2)_p-$, $-O(CH_2CH_2O)_q(CH_2)_p-$, $-O(CH_2)_p-$, $-N(R^1)(CH_2)_p-$, $-SO_2N(R^1)(CH_2)_pO(CH_2)_r-$, $-CON(R^1)(CH_2)_pO(CH_2)_r-$, $-OASO_2N(R^1)(CHR^1)_pOA-$ or $-(CH_2)_p(CHOH)_s(CH_2)_r-$, in which A represents an alkylene group or an arylene group, each R^1 and R^2 independently represent a hydrogen atom or an alkyl group having from 1 to 6 carbon atoms, p , r and s independently represent from 0 to 6 and q represents from 1 to 20.

5. A method according to any preceding claim, wherein the content ratio of the anionic fluorine-containing surfactant to the cationic fluorine-containing surfactant is from 1:10 to 10:1 in terms of mole ratio.

6. A method according to any preceding claim, wherein the total content of the anionic fluorine-containing surfactant and the cationic fluorine-containing surfactant is from 0.1 to 1000 mg/m².

7. A method according to any preceding claim, wherein the binder comprises gelatin.

8. A method according to claim 7, wherein the gelatin is an acid-processed gelatin or an amino group-inactivated gelatin.

9. A method according to claim 7 or 8, wherein the gelatin content of the ink receiving layer is from 3 to 50 g/m².

10. A method according to any preceding claim, wherein the binder comprises gelatin and a water-soluble polymer.

11. A method according to claim 10, wherein the water-soluble polymer is a polyvinyl pyrrolidone or a polyvinyl alcohol.

12. A method according to claim 10 or 11, wherein the content ratio of the water-soluble polymer to the gelatin is from

10 to 70 weight%.

13. A method of any preceding claim, wherein the support is a polyester film or a resin-covered paper.

5 Patentansprüche

1. Verfahren zur Tintenstrahlaufzeichnung bzw. Tintenstrahlaufzeichnungsverfahren, bei welchem eine Druckfarbe auf Wasserbasis auf ein Aufzeichnungsblatt ausgestoßen wird, wobei das Aufzeichnungsblatt einen Träger und eine darauf vorgesehene (Druck)farbenaufnahmeschicht, umfassend ein Bindemittel, ein anionisches, fluorhaltiges
10 Netzmittel und ein kationisches, fluorhaltiges Netzmittel umfaßt und wobei das anionische, fluorhaltige Netzmittel aus $C_{10}F_{21}(CH_2)_{10}COOH$ oder einer Verbindung der folgenden Formel (FA'):



15 worin bedeuten:

Rf eine fluorhaltige Alkylgruppe mit 2 bis 30 Kohlenstoffatomen oder eine Arylgruppe mit einer fluorhaltigen Alkylgruppe mit 2 bis 30 Kohlenstoffatomen;

D eine zweiwertige Gruppe mit mindestens einer verbindenden Gruppe aus -O-, -COO-, -CON(R₁)- und -
20 SO₂N(R₁)-sowie 1 bis 5 Kohlenstoffatom(en);

R₁ eine Alkylgruppe mit 1 bis 5 Kohlenstoffatom(en);

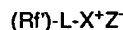
t = 1 oder 2;

Y -COOM-, -SO₃M oder -OSO₃M und

M ein Wasserstoffatom oder ein Alkalimetall oder ein quaternäres Ammoniumsalz,
25 besteht.

2. Verfahren nach Anspruch 1, wobei D für eine zweiwertige Gruppe mit einer -SO₂N(R₁)-Bindung steht und R₁ eine Alkylgruppe mit 1 bis 5 Kohlenstoffatom(en) darstellt.

- 30 3. Verfahren nach Anspruch 1 oder 2, wobei es sich bei dem kationischen, fluorhaltigen Netzmittel um eine Verbindung der folgenden Formel (FK):



35 worin bedeuten:

Rf' eine fluorhaltige Kohlenwasserstoffgruppe mit 1 bis 20 Kohlenstoffatom(en);

L eine chemische Bindung oder -SO₂N(R¹)(CH₂)_p-, -CON(R¹)(CH₂)_p-, -OASO₂N(R¹)(CH₂)_p-, -
40 OACON(R¹)(CH₂)_p-, -OAO(CH₂)_p-, -OA(CH₂)_p-, -O(CH₂CH₂O)_q(CH₂)_p-, -O(CH₂)_p-, -N(R¹)(CH₂)_p-, -
SO₂N(R¹)(CH₂)_pO(CH₂)_r-, -CON(R¹)(CH₂)_pO(CH₂)_r-, -OASO₂N(R¹)(CHR¹)_pOA- oder -
(CH₂)_p(CHOH)_s(CH₂)_r- mit A gleich einer Alkylgruppe oder einer Arylgruppe, jeder Rest R¹ und R² unab-
hängig voneinander gleich einem Wasserstoffatom oder einer Alkylgruppe mit 1 bis 6 Kohlenstoffatom(en), p,
r und s unabhängig voneinander gleich 0 bis 6 und q gleich 1 bis 20;

X⁺ ein Kation und

45 Z⁻ ein Anion,
handelt.

4. Verfahren nach Anspruch 3, wobei L aus -SO₂N(R¹)(CH₂)_p-, -CON(R¹)(CH₂)_p-, -OASO₂N(R¹)(CH₂)_p-, -
50 OACON(R¹)(CH₂)_p-, -OAO(CH₂)_p-, -OA(CH₂)_p-, -O(CH₂CH₂O)_q(CH₂)_p-, -O(CH₂)_p-, -N(R¹)(CH₂)_p-, -
SO₂N(R¹)(CH₂)_pO(CH₂)_r-, -CON(R¹)(CH₂)_pO(CH₂)_r-, -OASO₂N(R¹)(CHR¹)_pOA- oder -(CH₂)_p(CHOH)_s(CH₂)_r- mit
A gleich einer Alkylgruppe oder einer Arylgruppe, jeder Rest R¹ und R² unabhängig voneinander gleich einem
Wasserstoffatom oder einer Alkylgruppe mit 1 bis 6 Kohlenstoffatom(en), p, r und s unabhängig voneinander gleich
0 bis 6 und q gleich 1 bis 20 ausgewählt ist.

5. Verfahren nach einem der vorhergehenden Ansprüche, wobei das molare Anteilsverhältnis anionisches, fluorhaltiges Netzmittel/kationisches fluorhaltiges Netzmittel 1/10 bis 10/1 beträgt.

6. Verfahren nach einem der vorhergehenden Ansprüche, wobei der Gesamtgehalt an anionischem, fluorhaltigem

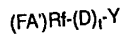
Netzmittel und kationischem, fluorhaltigem Netzmittel 0,1 - 1000 mg/m² beträgt.

7. Verfahren nach einem der vorhergehenden Ansprüche, wobei das Bindemittel Gelatine umfaßt.
8. Verfahren nach Anspruch 7, wobei es sich bei der Gelatine um eine säurebehandelte Gelatine oder eine Gelatine mit inaktivierten Aminogruppen handelt.
9. Verfahren nach Anspruch 7 oder 8, wobei der Gelatinegehalt der (Druck)farbenaufnahmeschicht 3 - 50 g/m² beträgt.
10. Verfahren nach einem der vorhergehenden Ansprüche, wobei das Bindemittel Gelatine und ein wasserlösliches Polymer umfaßt.
11. Verfahren nach Anspruch 10, wobei es sich bei dem wasserlöslichen Polymer um ein Polyvinylpyrrolidon oder einen Polyvinylalkohol handelt.
12. Verfahren nach Anspruch 10 oder 11, wobei der Anteil des wasserlöslichen Polymers, relativ zu Gelatine, 10 - 70 Gew.-% beträgt.
13. Verfahren nach einem der vorhergehenden Ansprüche, wobei es sich bei dem Träger um einen Polyesterfilm oder ein mit Harz beschichtetes bzw. kaschiertes Papier handelt.

Revendications

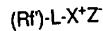
1. Procédé d'enregistrement à jet d'encre, comprenant l'étape consistant à :

envoyer des jets d'encre à base d'eau sur une feuille d'enregistrement, la feuille d'enregistrement comprenant un support et, sur celui-ci, une couche recevant l'encre comprenant un liant, un agent tensioactif fluoré anionique et un agent tensioactif fluoré cationique, l'agent tensioactif fluoré anionique étant C₁₀F₂₁(CH₂)₁₀COOH ou un composé de formule (FA') suivante :



dans laquelle Rf représente un groupe alkyle fluoré ayant de 2 à 30 atomes de carbone ou un groupe aryle ayant un groupe alkyle fluoré de 2 à 30 atomes de carbone ; D représente un groupe divalent ayant au moins un des groupes de liaison -O-, -COO-, -CON(R₁)- et -SO₂N(R₁)- et ayant de 1 à 5 atomes de carbone ; R₁ représente un groupe alkyle ayant de 1 à 5 atomes de carbone ; t représente 1 ou 2 ; Y représente -COOM-, -SO₃M ou -OSO₃M ; et M représente un atome d'hydrogène, un métal alcalin ou un sel d'ammonium quaternaire.

2. Procédé selon la revendication 1, dans lequel D est un groupe divalent ayant une liaison -SO₂N(R₁)- et R₁ est un groupe alkyle ayant de 1 à 5 atomes de carbone.
3. Procédé selon la revendication 1 ou 2, dans lequel l'agent tensioactif fluoré cationique est un composé de formule (FK) suivante :



dans laquelle Rf' représente un groupe hydrocarboné fluoré ayant de 1 à 20 atomes de carbone ; L représente une liaison chimique ou -SO₂N(R¹)(CH₂)_p-, -CON(R¹)(CH₂)_p-, -OASO₂N(R¹)(CH₂)_p-, -OACON(R¹)(CH₂)_p-, -OAO(CH₂)_p-, -OA(CH₂)_p-, -O(CH₂CH₂O)_q(CH₂)_p-, -O(CH₂)_p-, -N(R¹)(CH₂)_p-, -SO₂N(R¹)(CH₂)_pO(CH₂)_r-, -CON(R¹)(CH₂)_pO(CH₂)_r-, -OASO₂N(R¹)(CHR¹)_pOA- ou -(CH₂)_p(CHOH)_s(CH₂)_r-, où A représente un groupe alkylène ou un groupe arylène, chaque R¹ et R² représente indépendamment un atome d'hydrogène ou un groupe alkyle ayant de 1 à 6 atomes de carbone, p, r et s représentent indépendamment de 0 à 6 et q représente de 1 à 20 ; X⁺ représente un cation ; et Z⁻ représente un anion.

4. Procédé selon la revendication 3, dans lequel L est choisi parmi -SO₂N(R¹)(CH₂)_p-, -CON(R¹)(CH₂)_p-, -OASO₂N(R¹)(CH₂)_p-, -OACON(R¹)(CH₂)_p-, -OAO(CH₂)_p-, -OA(CH₂)_p-, -O(CH₂CH₂O)_q(CH₂)_p-, -O(CH₂)_p-,

$N(R^1)(CH_2)_p-$, $-SO_2N(R^1)(CH_2)_pO(CH_2)_r-$, $-CON(R^1)(CH_2)_pO(CH_2)_r-$, $-OASO_2N(R^1)(CHR^1)_pOA-$ ou $-(CH_2)_p(CHOH)_s(CH_2)_r-$, où A représente un groupe alkylène ou un groupe arylène, chaque R^1 et R^2 représente indépendamment un atome d'hydrogène ou un groupe alkyle ayant de 1 à 6 atomes de carbone, p, r et s représentent indépendamment de 0 à 6 et q représente de 1 à 20.

- 5 5. Procédé selon l'une quelconque des revendications précédentes, dans lequel le rapport de l'agent tensioactif fluoré anionique à l'agent tensioactif fluoré cationique est de 1:10 à 10:1 en rapport molaire.
- 10 6. Procédé selon l'une quelconque des revendications précédentes, dans lequel la proportion totale de l'agent tensioactif fluoré anionique et de l'agent tensioactif fluoré cationique est de 0,1 à 1000 mg/m².
7. Procédé selon l'une quelconque des revendications précédentes, dans lequel le liant comprend de la gélatine.
- 15 8. Procédé selon la revendication 7, dans lequel la gélatine est une gélatine traitée par un acide ou une gélatine à groupe amino inactivé.
9. Procédé selon la revendication 7 ou 8, dans lequel la proportion de gélatine de la couche recevant l'encre est de 3 à 50 g/m².
- 20 10. Procédé selon l'une quelconque des revendications précédentes, dans lequel le liant comprend de la gélatine et un polymère hydrosoluble.
11. Procédé selon la revendication 10, dans lequel le polymère hydrosoluble est une polyvinylpyrrolidone ou un alcool polyvinylique.
- 25 12. Procédé selon la revendication 10 ou 11, dans lequel la proportion du polymère hydrosoluble par rapport à la gélatine est de 10 à 70 % en poids.
- 30 13. Procédé selon l'une quelconque des revendications précédentes, dans lequel le support est un film polyester ou un papier revêtu de résine.